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<b>(21) International Application Number: PCT/US94/09048</b> <b>(22) International Filing Date: 10 August 1994 (10.08.94)</b>  <b>(30) Priority Data:</b> 08/106,434      13 August 1993 (13.08.93)      US  <b>(71) Applicant: ZONAGEN, INC. [US/US]; 2408 Timberloch</b> Place, B-4, The Woodlands, TX 77380 (US).  <b>(72) Inventors: ZORNIOTTI, Adrian, W.; 33 E. 74th Street, New</b> York, NY 10021 (US). GIOCO, Diane-Marie; 199 First Avenue, West Haven, CT 06516 (US).  <b>(74) Agent: CLOUGH, David, W.; Marshall, O'Toole, Gerstein,</b> Murray & Borun, 6300 Sears Tower, 233 S. Wacker Drive, Chicago, IL 60606-6402 (US).		<b>(81) Designated States: AU, BR, CA, CN, ES, JP, KR, NO, NZ,</b> European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title: METHODS FOR MODULATING THE HUMAN SEXUAL RESPONSE</b>  <b>(57) Abstract</b>  The invention is directed to improved methods for modulating the human sexual response by administering a vasodilator to the circulation by a route of administration that avoids the "first-pass" effect.		

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## METHODS FOR MODULATING THE HUMAN SEXUAL RESPONSE

### CROSS REFERENCE TO RELATED APPLICATIONS

This application is a continuation-in-part of co-pending  
5 commonly-assigned U.S. Patent Application Serial No. 08/106,434, filed  
August 13, 1993.

### FIELD OF THE INVENTION

The application is directed to improved methods for modulating  
the human sexual response by administering vasodilator agents to the  
10 circulation of a human via transmucosal, transdermal, intranasal or rectal  
routes of administration that obviate "first-pass" deleterious effects on such  
agents.

### BACKGROUND OF THE INVENTION

The human sexual response in both males and females results  
15 from a complex interplay of psychological, hormonal, and other physiological  
influences. One important aspect of human sexual response that is common  
to both men and women is the erectile response which itself results from an  
interplay between the autonomic nervous system, the endocrine system, and  
the circulatory system.

20 Failure of the erectile response is most common in men and is  
referred to as impotence. Impotence is the inability of a male to achieve or  
sustain a penile erection sufficient for vaginal penetration and intercourse.  
Numerous approaches have been taken in attempts to treat impotence. These  
approaches include the use of external or internally implanted penile  
25 prosthesis. (See, *e.g.*, U.S. Patent No. 5,065,744, to Zumanowsky). A  
variety of drugs and methods for administering drugs have also been used in

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attempts to treat impotence. For example, U.S. Patent No. 3,943,246 to Stürmer addresses treatment of impotence in men by buccal and peroral administration of daily doses of 300-1500 international units (I.U.) of oxytocin or daily divided doses of 150-250 I.U. of desamino-oxytocin. The patent  
5 states that the buccal administration of 100 I.U. three times a day for 14 days results in improvement of *impotentia erectionis* in 12 of the 16 patients treated.

U.S. Patent No. 4,530,920 to Nestor *et al.* suggests the possibility that administration of nonapeptide and decapeptide analogs of  
10 luteinizing hormone releasing hormone agonists may be useful in the induction or enhancement of sexual behavior or therapy for impotence or frigidity. Nestor *et al.* suggest numerous routes of administration of the analogs including buccal, sublingual, oral, parenteral (including subcutaneous, intramuscular, and intravenous administration), rectal, vaginal, and others.

15 U.S. Patent No. 4,139,617 to Grunwell *et al.* suggests buccal and other routes of administration of 19-oxygenated-androst-5-enes for the endocrine mediated enhancement of the libido in humans.

U.S. Patent No. 4,863,911 to Anderson *et al.* addresses methods for treating sexual dysfunction in mammals using a biooxidizable,  
20 blood-brain barrier penetrating estrogen derivative. One of the purported objects of the Anderson *et al.* invention is the treatment of "psychological impotence" in males. Test results showed that the drugs used in the study stimulated mounting behavior, intromission, and mount latency in castrated rats.

25 A number of publications have proposed the use of various vasodilators for the treatment of impotence in males. Attempts to utilize vasodilators for the treatment of impotence were prompted because a significant percentage of cases of impotence were noted to be vasculogenic, i.e., resulting from vascular insufficiency.

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Voss *et al.*, U.S. Patent No. 4,801,587, issued January 31, 1989, addresses the use of an ointment containing a vasodilator and a carrier agent for topical application to the penis of impotent men. The Voss *et al.* patent also describes application of such an ointment into the urethra of the penis using a catheter as well as a multi-step regimen for applying a vasodilator to the skin of the penis. In addition, Voss *et al.* proposes the surgical removal of a portion of the fibrous sheath surrounding the corpora cavernosum, thereby facilitating the penetration of a vasodilator-containing ointment into the corpora cavernosum. Vasodilators suggested for use by Voss *et al.* include papaverine, hydralazine, sodium nitroprusside, phenoxybenzamine, and phentolamine. The Voss *et al.* patent, however, provides no information regarding the actual efficacy of the treatments proposed or the nature of the response to such treatments.

U.S. Patent No. 4,127,118 to Latorre describes treating male impotence by directly injecting the vasodilating drugs into the corpus cavernosum and the corpus spongiosum of the penis using a syringe and one or more hypodermic needles. More particularly, the Latorre patent proposes the intracavernosal and intraspongiosal injection of sympathomimetic amines such as nylidrin hydrochloride, adrenergic blocking agents such as tolazoline hydrochloride, and direct acting vasodilators such as isoxsuprine hydrochloride and nicotiny alcohol.

Brindley, G.S. (*Br. J. Pharmac.* 87:495-500 1986) showed that, when injected directly into the corpus cavernosum using a hypodermic needle, certain smooth muscle relaxing drugs including phenoxybenzamine, phentolamine, thymoxamine, imipramine, verapamil, papaverine, and naftidrofuryl caused erection. This study noted that injection of an "appropriate dose of phenoxybenzamine or papaverine is followed by an unrelenting erection lasting for hours." Injection of the other drugs studied induced erections lasting from about 11 minutes to about 6.5 hours. Zorogniotti *et al.*, *J. Urol.* 133:39-41 (1985) demonstrated that the

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intracavernosal injection of a combination of papaverine and phentolamine could result in an erection in otherwise impotent men. Similarly, Althof *et al.* *J. Sex Marital Ther.* 17(2): 101-112 (1991) reported that intracavernosal injection of papaverine hydrochloride and phentolamine mesylate resulted in improved erectile ability in about 84% of patients injected. However, in that study the dropout rate was 57%, fibrotic nodules developed in 26% of the patients, 30% of the patients developed abnormal liver function values, and bruising occurred in 19% of the patients.

Other studies describing intracavernosal injection of drugs using hypodermic needles for the treatment of impotence include: Brindley, *J. Physiol.* 342:24P (1983); Brindley, *Br. J. Psychiatr.* 143:312-337 (1983); Virag, *Lancet* ii:978 (1982); and Virag, *et al.*, *Angiology* 35:79-87 (1984).

While intracavernosal injection may be useful for inducing erections in impotent men, the technique has numerous drawbacks. Obvious drawbacks include pain, risk of infection, inconvenience and interference with the spontaneity of the sex act. Priapism (prolonged and other painful erection) also appears to be a potential problem when using injection methods. See, *e.g.* Brindley, (1986). Another problem arising in some cases of intracavernosal injection involves the formation of fibrotic lesions in the penis. See, *e.g.*, Corriere, *et al.*, *J. Urol.* 140:615-617 (1988) and Larsen, *et al.*, *J. Urol.* 137:292-293 (1987).

Phentolamine, which has been shown to have the potential to induce erection when injected intracavernosally, has also been the subject of oral administration to test its effects in men having non-specific erectile insufficiency (Gwinup, *Ann. Int. Med.* 15 July 1988, pp. 162-163). In that study, 16 patients ingested either a placebo or a 50 mg orally administered dose of phentolamine. Eleven of the 16 patients (including three placebo-treated patients) became tumescent, became more responsive to sexual stimulation, and were able to achieve an erection sufficient for vaginal penetration after waiting 1.5 hours to attempt intercourse.

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Sonda *et al.* *J. Sex & Marital Ther.* 16(1): 15-21 (year) reported that yohimbine ingestion resulted in subjective improvement in erectile ability in 38% of impotent men treated, but only 5% of the treated patients reported complete satisfaction.

5           Of interest to the background of the invention are the disclosures of Stanley *et al.*, U.S. Patent No. 4,885,173, which addresses methods for non-invasively administering drugs having cardiovascular or renal vascular activity through use of a lollipop assertedly facilitating drug absorption through the mucosal tissues of the mouth, pharynx, and esophagus.  
10   The Stanley *et al.* patent proposes that a large number of lollipop-administered drugs may improve cardiovascular function including drugs exhibiting direct vasodilating effects, calcium channel blockers,  $\beta$ -adrenergic blocking agents, serotonin receptor blocking agents, angina blocking agents, other anti-hypertensive agents, cardiac stimulating agents, and agents which improve  
15   renal vascular function.

U.S. Patent No. 5,059,603 to Rubin describes the topical administration to the penis of isoxsuprine and caffeine, and nitroglycerine and caffeine along with suitable carrier compounds for the treatment of impotence.

20           There continues to exist a need in the art for effective means for modulating human sexual response and especially for enhancing erectile ability in males suffering from impotence. Ideally, such means would be convenient and simple to use, would not require a constant dosage regimen or even multiple doses to achieve desired results, would be non-invasive and would allow a rapid and predictable capacity for onset of erectile function on  
25   demand and in response to normal sexual stimulation.

#### SUMMARY OF THE INVENTION

The present invention provides improved methods for modulating the human sexual response in a human by administering a vasodilator agent to the circulation in an amount effective to increase blood

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flow to the genitalia. According to the invention, modulation of male and female human sexual response is provided on demand by administering an effective vasodilating amount of the agent by a route selected from the group consisting of transmucosal, transdermal, intranasal and rectal. Vasodilating agents useful in the present invention include, but are not limited to, the group consisting of phentolamine mesylate, phentolamine hydrochloride, phenoxybenzamine, yohimbine, organic nitrates (e.g. nitroglycerin), thymoxamine, imipramine, verapamil, isoxsuprine, naftidrofuryl, tolazoline and papaverine. The presently preferred agent is phentolamine mesylate. The presently preferred administrative route is transmucosal, especially buccal.

The present invention is specifically directed to improved methods for treating male impotence by administering a vasodilator agent in an amount effective to increase blood flow to the penis wherein erectile ability on demand is permitted by administering the agent by an administrative route selected from the group consisting of transmucosal, transdermal, intranasal, and rectal.

Preferably, the amount of vasodilating agent used in the practice of the invention for treatment of male impotence is effective to improve erectile ability in from about 1 minute to about 60 minutes following administration of the agent.

The invention is also specifically directed to methods for modulating the excitation and plateau phases of the female sexual response on demand by transmucosal, transdermal, intranasal, or rectal administration of an effective amount of vasodilator agent.

The methods of the present invention are also useful in preparation for sexual intercourse by virtue of the ability to modulate the sexual response in both males and females.

The present invention is also directed to the use of a drug having vasodilator activity for the manufacture of a medicament for transmucosal, transdermal, intranasal, and rectal administration to modulate

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sexual response in a human. Vasodilator drugs useful for manufacturing the medicament include, but are not limited to, phentolamine mesylate, phentolamine hydrochloride, phenoxybenzamine yohimbine, organic nitrates, thymoxamine, imipramine, verapamil, isoxsuprine, naftidrofuryl, tolazoline, and papaverine.

Numerous other advantages of the present invention will be apparent from the following detailed description of the invention including the accompanying examples and the appended claims.

#### **BRIEF DESCRIPTION OF THE DRAWINGS**

Fig. 1 is a graph depicting the percentage change in penile arterial blood velocity after buccal administration of a 20 mg dose of phentolamine mesylate.

#### **DETAILED DESCRIPTION**

The human sexual response in both the male and female involves a complex interplay between endocrine, neurological and psychological components which result in certain physiological and anatomical responses in both men and women.

While there are obvious differences in the sexual response between men and women, one common aspect of the sexual response is the erectile response. The erectile response in both males and females is result of engorgement of the erectile tissues of the genitalia with blood in response to sexual stimulation (physical, psychological, or both).

The vasculature which serves erectile tissue in both men and women is similar. In particular, in both men and women, the arterial circulation to the erectile tissues of the genitalia derives from the common iliac artery which branches from abdominal aorta. The common iliac artery bifurcates into the internal and external iliac arteries. The internal pudic artery arises from the smaller of two terminal branches of the anterior trunk

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of the internal iliac artery. In the female, the internal pudic artery branches into the superficial perineal artery which supplies the labia pudenda. The internal pudic artery also branches into the artery of the bulb which supplies the bulbi vestibuli and the erectile tissue of the vagina. The artery of the corpus cavernosum, another branch of the internal pudic artery supplies the cavernous body of the clitoris. Still another branch of the internal pudic artery is the arteria dorsalis clitoridis which supplies the dorsum of the clitoris and terminates in the glans and membranous folds surrounding the clitoris which correspond to the prepuce of the male.

In the male, the internal pudic artery branches into the dorsal artery of the penis (which itself branches into a left and right branch) and the artery of the corpus cavernosum, all of which supply blood to the corpus cavernosum. The dorsal artery of the penis is analogous to the artery dorsalis clitoridis in the female, while the artery of the corpus cavernosum in the male is analogous to the artery of the same name in the female.

The male erectile response is regulated by the autonomic nervous system which controls blood flow to the penis via the interaction of peripheral nerves associated with the arterial vessels in and around the corpus cavernosum. In the non-aroused or non-erect state, the arteries serving the corpus cavernosum are maintained in a relatively constricted state, thereby limiting the blood flow to the corpus cavernosum. However, in the aroused state, the smooth muscles associated with the arteries relax under the influence of catecholamines and blood flow to the corpus cavernosum greatly increases, causing expansion and rigidity of the penis. Brindley, *supra* (1986) hypothesizes that smooth muscle contraction opens valves through which blood can flow from the corpus cavernosum into the extracavernosal veins. According to Brindley (1986), when the relevant smooth muscles relax, the valves close diminishing venous outflow from the corpus cavernosum. When accompanied by increased arterial blood flow into the corpus cavernosum, this results in engorgement of the corpus cavernosum and an erection.

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The pre-orgasmic sexual response in females can be broken down into distinct phases. Both the excitement phase and the plateau phase involve vasodilation and engorgement (vasocongestion) of the genitalia with arterial blood in a manner analogous to the male erectile response.

5           The excitement phase of the female sexual response is characterized by vasocongestion in the walls of the vagina which leads to the transudation of vaginal fluids and vaginal lubrication. Further, the inner one-third of the vaginal barrel expands and the cervix and the body of the uterus become elevated. This is accompanied by the flattening and elevation of the  
10   labia majora and an increase in clitoral size. [Kolodny *et al.*, *Textbook of Sexual Medicine*, Little and Brown, Boston, MA (1979)].

          The plateau phase follows the excitement phase in the female sexual response and is characterized by prominent vasocongestion in the outer one-third of the vagina, causing a narrowing of the opening of the vagina and  
15   a retraction of the shaft and the glans of the clitoris against the symphysis pubis. These responses are also accompanied by a marked vasocongestion of the labia. [Kolodny, *supra* (1979)].

          The vasocongestive aspects of the female sexual response are not restricted to the genitalia in that areolar engorgement also occurs,  
20   sometimes to the extent that it masks the antecedent nipple erection that usually accompanies the excitement phase.

          The failure of the erectile response in men to the extent that vaginal penetration and sexual intercourse cannot be achieved is termed impotence. Impotence has numerous possible causes which can be broken  
25   down into several general classifications. Endocrine related impotence can result from primary gonadal failure, advanced diabetes mellitus, hypothyroidism, and as one of the secondary sequelae of pituitary adenoma, idiopathic or acquired hypogonadism, hyperprolactinemia and other endocrine abnormalities.

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Chronic systemic illnesses such as cirrhosis, chronic renal failure, malignancies and other systemic diseases can also cause impotence. Neurogenic impotence arising in the central nervous system can be caused by temporal lobe disorders caused by trauma, epilepsy, neoplasms and stroke, intramedullary spinal lesions, paraplegia, and demyelinating disorders. Neurogenic causes of impotence arising in the peripheral nervous system include somatic or autonomic neuropathies, pelvic neoplasms, granulomas, trauma, and others. Urologic causes of impotence include complete prostatectomy, local trauma, neoplasms, Peyronie's disease, and others.

As many as half the cases of male impotence may be psychogenic because there is no readily-ascertainable organic cause for the disorder. Even when there appears to be an underlying organic cause of impotence, psychologic factors may play a role in the disorder.

The present invention is designed to modify the circulatory aspects of the erectile response using vasoactive agents administered to the circulation by a route selected from the group consisting of transmucosal, transdermal, intranasal, and rectal.

A number of vasoactive agents may be used in the practice of the present invention based on demonstrated systemic efficacy as vasodilators. Useful vasodilating drugs include those generally classified as  $\alpha$ -adrenergic antagonists, sympathomimetic amines and those agents which exhibit direct relaxation of vascular smooth muscle. Exemplary  $\alpha$ -adrenergic antagonists include phentolamine hydrochloride, phentolamine mesylate, phenoxybenzamine, tolazoline, dibenamine, yohimbine, and others. Phentolamine mesylate is a preferred  $\alpha$ -adrenergic agent vasodilator for use preferred practice of the present invention. An exemplary sympathomimetic amine contemplated for use in the method of the present invention is nylidrin and use of other sympathomimetic amines having vasodilating activity is also contemplated.

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Nicotinic acid (or nicotiny alcohol) has a direct vasodilating activity useful in the practice of the present invention. Also contemplated is the use of papaverine, a non-specific smooth muscle relaxant which possesses vasodilating activity and which has been used to treat male impotence by direct injection into the corpus cavernosum either alone or in combination with other drugs such as phentolamine. Organic nitrates such as nitroglycerine and amyl nitrate have pronounced vasodilating activity by virtue of their ability to relax vascular smooth muscle and are thus contemplated for use according to the invention. Other vasoactive drugs useful in the practice of the present invention include, without limitation, thymoxamine, imipramine, verapamil, naftidrofuryl, and isoxsuprine.

In the practice of the present invention, vasoactive agents are administered by the transmucosal, intranasal, transdermal, or rectal routes of administration such that the agent is conveyed in circulation to the site of action prior to entering portal circulation, thereby avoiding the "first-pass" effect described below.

Oral administration of a drug in an attempt to effect delivery to a specific site within circulation may have several drawbacks. First, drug absorption is limited by gastrointestinal transit time and thus, the rapidity of onset of drug action may be limited. Second, the drug may be inactivated by (e.g. by hydrolysis) in the low pH environment of the stomach and or by other chemical or biochemical interactions in the intestines. Third, when an orally ingested drug reaches the intestine, it is absorbed into the portal circulation and delivered to the liver where it can be metabolized and inactivated. Hepatic inactivation following absorption of a drug from the gastrointestinal tract is referred to as "first-pass" effect or "presystemic" elimination and, along with poor absorption and slow transit times through the gastrointestinal tract, functions to require larger oral doses of drugs than may be necessary with other routes of administration. This, in turn, can account for delays in the onset of the therapeutic effect of a drug. [Goodman *et al.*, *The*

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*Pharmacological Basis of Therapeutics*, Macmillan Publishing Co., New York, N.Y. (1975)].

Transmucosal, transdermal, intranasal, and rectal routes of administration of vasodilators according to the present invention avoids the first-pass effect allowing administration of the vasodilator a short time prior to the projected initiation of intercourse ("on demand") and eliminating the need for repeated administration of the drug. Methods of the present invention are thus more convenient and help minimize any side-effects that can arise as a result of continuous or daily administration of the drugs. In addition, methods of the present invention allow more spontaneity in sexual activity than allowed by other methods such as intracavernosal injection of vasodilators.

Formulations for effecting transmucosal delivery of vasodilators according to the present invention are well known in the art. For purposes of the present invention, "transmucosal delivery" generally refers to delivery of the drug to the oral or pharyngeal mucosa and includes buccal delivery, sublingual delivery, and delivery to the pharyngeal mucosa, but not to the stomach. Buccal delivery may be accomplished by preparing a tablet or lozenge comprising, for example, compressed lactose and an effective amount of one or more vasodilators. Other suitable tablet compositions include, but are not limited to, a combination of an effective dose of a vasodilator, and carrier substances, tablet-binding compounds and flavoring agents such as those described in U.S. Patent No. 3,943,246 to Stürmer, which is incorporated herein by reference. Vasoactive agents may also be compounded with a variety of pharmaceutical excipients including binders such as gelatin and/or corn starch or pharmaceutically acceptable gums such as gum tragacanth. Vasoactive agents may also be combined in a hard candy (which may be dissolved in the mouth) or in a chewing gum, to provide buccal or sublingual delivery to the oral mucosa.

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Vasodilators may also be administered transmucosally by impregnating a filter paper strip or a filter paper disc with an effective amount of a vasodilator. The filter paper strip or disc may then be placed between the cheek and gum (buccally) for delivery to the vasculature of the genitalia  
5 without encountering first-pass effects. Other transmucosal delivery systems such as lollipops (as described in U.S. Patent No. 4,885,173 to Stanley) are well known in the art are expected to be useful in the practice of the present invention.

Transdermal administration of vasodilators according to the  
10 present invention also eliminates the "first-pass" effect occurring when drugs are administered orally or into circulatory flow leading to the liver.

Transdermal delivery systems are well known in the art and involve what are sometimes referred to as transdermal "patches". Exemplary transdermal patches typically comprise: (1) a impermeable backing layer  
15 which may be made up of any of a wide variety of plastics or resins, *e.g.* aluminized polyester or polyester alone or other impermeable films; and (2) a drug reservoir comprising, for example, a vasodilator in combination with mineral oil, polyisobutylene, and alcohols gelled with USP hydroxymethylcellulose. Other exemplary drug reservoir layers may  
20 comprise, for example, acrylic-based polymer adhesives with resinous cross-linking agents which provide for diffusion of the drug from the reservoir to the surface of the skin. The transdermal patch may also have a delivery rate-controlling membrane such as a microporous polypropylene disposed between the reservoir and the skin. Ethylene-vinyl acetate copolymers and other  
25 microporous membranes may also be used. Typically, an adhesive layer is provided which may comprise an adhesive formulation such as mineral oil and polyisobutylene combined with the vasoactive agent.

Other typical transdermal patches may comprise three layers:  
(1) an outer layer comprising a laminated polyester film; (2) a middle layer  
30 containing a rate-controlling adhesive, a structural non-woven material and the

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vasodilator; and (3) a disposable liner that must be removed prior to use. Transdermal delivery systems may also involve incorporation of highly lipid soluble carrier compounds such as dimethyl sulfoxide (DMSO), to facilitate penetration of the skin. Other exemplary carriers include lanolin and glycerin.

5               Vasodilating drugs for use in transmucosal, transdermal, intranasal, or rectal delivery may be chemically modified by methods well known in the art to improve their lipid solubility and thus their ability to penetrate skin or mucosal surfaces.

              Rectal suppositories are also useful in the practice of the present  
10   invention for administration by a route which by-passes the "first-pass" effect. Exemplary suppositories comprise a vasodilating drug combined with glycerin, glycerol monopalmitate, glycerol, monostearate, hydrogenated palm kernel oil and fatty acids. Another exemplary suppository formulation includes ascorbyl palmitate, silicon dioxide, white wax, and cocoa butter in combination with  
15   an effective amount of a vasodilating drugs.

              The present invention is also directed to the use of nasal sprays for the administration of the vasodilators. Exemplary nasal spray formulations comprise a solution of vasodilating drug in physiologic saline or other pharmaceutically suitable carrier liquids. Nasal spray compression pumps are  
20   also well known in the art and can be calibrated to deliver a predetermined dose of the vasodilator solution.

              The examples set forth below are intended to be illustrative of the present invention and are not intended to limit the scope of the invention as set out in the appended claims.

25               Example 1 describes the effect of transmucosally administered phentolamine mesylate on penile arterial velocity. Examples 2 and 3 describe the effect of buccally administered phentolamine mesylate on erectile ability in impotent men. Example 4 describes the use of a variety of vasodilators in the practice of the present invention. Example 5 addresses practice of the  
30   present invention in modulating erectile response in females.

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### EXAMPLE 1

#### EFFECT OF TRANSMUCOSAL ADMINISTRATION OF PHENTOLAMINE MESYLATE ON PENILE ARTERIAL VELOCITY

5                   In order to assess whether or not buccally administered  
phentolamine mesylate could alter arterial blood flow in the penis, penile  
arterial velocity was measured close to the base of the right branch of the  
dorsal artery of the penis by doppler ultrasound velocimetry using a Dasonics  
400 DRF (Dasonics, Milpitas, CA). Settings, incidence of doppler beam,  
10                   and angle correction were maintained to achieved maximum velocity readings  
on each subject. The dorsal artery was selected for measurement because it  
was more accessible than the artery of the corpus cavernosum.

                  Velocities were measured before the transmucosal (buccal)  
administration of phentolamine mesylate (20 mg) and at 5, 15, 45 and 60  
15                   minutes after administration of the drug. Mean initial velocity was 10.4  
cm/sec. The data shown in Figure 1 shows the percentage increase in penile  
artery velocity versus time after administration of phentolamine mesylate and  
represent the mean of triplicate readings in six impotent subjects.

                  The results show that, within five minutes of placing the tablet  
20                   between the cheek and gum, arterial velocity rose by more than 50% above  
base line velocity. Within 25 minutes, arterial velocity peaked at more than  
100% above base line velocity after which velocity began to fall, reaching  
pretreatment levels after 1 hour. Buccal administration of 20 mg of  
phentolamine mesylate is thus shown to provide a suitable means for rapidly  
25                   altering penile arterial blood flow.

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## EXAMPLE 2

### EFFECT OF TRANSMUCOSAL (BUCCAL) ADMINISTRATION OF PHENTOLAMINE MESYLATE ON MALE ERECTILE ABILITY

5 The fact that buccal administration of 20 mg of phentolamine mesylate as in Example 1 resulted in a rapid increase in penile artery velocity supported utility of this route of administration of vasodilators in improving erectile ability in impotent men. Thus, studies were therefore conducted to determine the effect of transmucosal administration of phentolamine mesylate on erectile ability in impotent men.

10 All patients included in the study complained of an erectile dysfunction which either prevented vaginal penetration or was characterized by an inability to maintain an erection without ejaculation upon initiation of vaginal penetration. The duration of impotence was within a range of 0.5 years to 35 years in this patient population and the mean duration was 3.4  
15 years.

Prior to administration of phentolamine mesylate or placebo, a medical history was taken, a genital examination was performed, and penile vascular status was determined. Vascular status was determined by measuring brachial and penile systolic closing pressures and calculating a penile brachial  
20 index (PBI). PBI was calculated by dividing the penile systolic pressure by the brachial systolic pressure. Plethysmographic crest times (CT) were measured with a Penilab IV plethysmograph (Parkes, Aloha WA). Crest time is the time in seconds from the trough of the penile blood pressure curve to the next peak. The normal range of crest times is from about 1 second to about 1.8 seconds. A PBI of  $>0.9$  is considered normal while a PBI of  $<0.6$   
25 indicates vascular insufficiency.

For the purposes of this study, the patients were identified as having normal vascularity if both PBI and CT were in the normal range. Patients having both CT and PBI in the abnormal range were identified as

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having marked vascular insufficiency. Patients having one parameter in the abnormal range were identified as having moderate vascular insufficiency.

Age and diagnosis were generally not a factor in admission to the study, although patients with extreme age or severe penile vascular insufficiency were excluded. Patients having vascular or non-specific causes of impotence were admitted to the trial, as were patients having diabetes mellitus. Mean age of patients was 57.5 years (range: 25-74 years).

In order to test the effort of phentolamine mesylate on erectile ability, a single blind trial was conducted. Each patient was given two tablets; a placebo tablet comprising lactose alone, and a second lactose tablet containing 20 mg of phentolamine mesylate.

Patients were asked to place one tablet between the cheek and gum (buccal) 10-20 minutes before attempting coitus. Buccal administration was used as a paradigm of transmucosal delivery which, like all routes of delivery useful in the practice of the present invention, results in the drug reaching the vasculature of the target (genitalia) prior to passing through the liver. One or more days after using the first tablet, patients repeated the process with the second tablet. Patients were advised not to swallow the tablets or consume alcohol prior to using the tablets and were told not to expect erection without sexual stimulation.

Patients were told that either tablet might prove beneficial and were told to report results in terms of erection and vaginal penetration, partial erection, or failure to maintain an erection sufficient to permit vaginal penetration. Patients were also asked to report side effects. Results of this study are illustrated in Table 1 which shows the age, vascular status and effect of phentolamine mesylate on erectile ability in impotent men. In the right-hand columns of Table 1, the number 1 indicates a report of erection and vaginal penetration, 2 indicates reported failure to achieve erection, and 3 indicates report of a partial erection.

**TABLE 1**  
**VASCULAR STATUS AND THE EFFECTS OF BUCCAL ADMINISTRATION**  
**OF PHENTOLAMINE ON PENILE ERECTILE ABILITY IN IMPOTENT MEN**

Patient Number	Age	Crest Time	Penile Brachial Indices		Phentolamine (20 mg)	Placebo
			Right	Left		
1	60	1.5	0.75	0.87	3	2
2	60	2.4	0.79	0.82	1	2
3	64	2.4	0.78	0.81	1	2
4	61	2.4	0.72	0.72	2	2
5	60	2.2	0.84	0.82	1	2
6	39	1.4	0.77	0.79	1	1
7	46	1.6	0.91	1.00	2	2
8	33	1.4	0.76	0.78	3	2
9	44	1.6	0.99	0.89	2	2
10	36	2.4	0.90	0.90	2	2
11	66	2.2	0.91	0.90	1	2
12	45	1.8	0.67	0.87	1	1

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TABLE 1 cont'd.

Patient Number	Age	Crest Time	Penile Brachial Indices		Phentolamine (20 mg)	Placebo
			Right	Left		
13	50	2.0	0.81	0.74	2	2
14	62	1.6	0.82	0.97	3	2
15	51	1.8	0.78	0.89	3	2
16	44	2.6	0.69	0.69	1	1
17	38	2.0	1.00	0.87	2	2
18	49	1.4	0.93	0.94	1	2
19	49	2.2	0.89	1.04	2	2
20	53	2.4	0.80	0.86	2	2
21	65	1.6	1.05	0.96	1	2
22	54	1.6	0.80	0.81	2	1
23	65	1.8	0.95	0.93	1	1
24	63	1.6	1.00	0.99	2	1
25	64	2.0	0.86	0.77	3	2
26	54	1.2	0.92	0.88	1	2

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TABLE 1 cont'd.

Patient Number	Age	Crest Time	Penile Brachial Indices		Phentolamine (20 mg)	Placebo
			Right	Left		
27	65	2.6	0.98	0.93	2	2
28	64	2.8	0.90	0.80	2	2
29	67	2.4	0.84	0.89	2	2
30	68	2.2	0.74	0.71	1	2
31	60	2.0	1.03	1.00	2	2
32	51	1.6	0.79	0.75	2	2
33	72	2.8	1.05	1.07	1	1
34	59	2.4	0.55	0.56	1	2
35	56	1.6	0.68	0.79	1	2
36	59	2.4	0.85	0.79	2	2
37	58	3.2	0.65	0.53	2	1
38	50	2.2	0.89	0.98	2	2
39	66	2.4	0.59	0.54	1	3
40	51	2.8	0.66	0.55	3	2

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TABLE 1 cont'd.

Patient Number	Age	Crest Time	Penile Brachial Indices		Phentolamine (20 mg)	Placebo
			Right	Left		
41	53	2.8	0.69	0.78	2	2
42	62	2.2	0.93	0.95	2	2
43	66	2.0	0.80	0.82	1	1
44	56	2.0	0.65	0.73	3	2
45	74	2.9	0.86	0.87	2	2
46	63	2.2	1.10	1.08	2	2
47	72	2.0	0.55	0.67	2	2
48	70	2.4	0.81	0.83	2	2
49	69	2.4	0.95	0.95	2	2
50	55	2.0	0.82	0.80	3	2
51	55	2.4	1.01	0.92	2	3
52	65	2.4	0.81	0.80	3	2
53	60	2.0	0.74	0.83	2	2
54	62	1.6	0.87	0.88	1	2

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TABLE 1 cont'd.

Patient Number	Age	Crest Time	Penile Brachial Indices		Phentolamine (20 mg)	Placebo
			Right	Left		
55	53	2.2	1.02	0.98	2	2
56	37	1.6	0.96	0.87	1	2
57	55	2.0	0.71	0.56	1	2
58	25	1.8	0.89	0.91	2	2
59	51	2.0	1.02	1.14	2	2
60	35	2.0	0.85	0.98	2	2
61	69	2.8	0.93	0.97	2	2
62	40	2.3	0.95	0.91	2	2
63	54	2.0	0.98	0.86	1	2
64	76	2.2	0.70	0.71	3	2
65	53	2.4	0.92	0.90	2	2
66	59	2.6	0.79	0.69	2	2
67	59	2.0	0.83	0.87	2	2
68	53	3.2	0.58	0.78	1	2

TABLE 1 cont'd.

Patient Number	Age	Crest Time	Penile Brachial Indices		Phentolamine (20 mg)	Placebo
69	38	2.2	Right	0.88	0.85	2
			Left			2

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The data in Table 1 illustrates that buccal administration of 20 mg of phentolamine mesylate resulted in improved erectile ability within 10 minutes to 20 minutes after buccal administration of the drug. The response was characterized by improved erectile ability upon sexual stimulation and thus, the response closely mimicked the normal sexual response in men. The fact that the effect of the drug was seen within 10-20 minutes after a single administration can be characterized as a response occurring "on demand" in that multiple doses and/or a long waiting time before onset of improved erectile ability were not required. The rapid increase in penile artery velocity within five minutes of administration of phentolamine mesylate (as indicated in Example 1) suggests that improve erectile ability may actually occur sooner than 10 minutes after administration. The "on demand" aspect of the method of the present invention allows a more natural and more spontaneous approach to intercourse and eliminates the need for multiple doses, and thus, reduces the frequency of undesirable side effects. Although a 20 mg dose of phentolamine mesylate was used in the present study, doses from about 5 mg to about 80 mg of phentolamine mesylate are within the scope of the present invention as individual responsiveness to the drug may vary *e.g.*, on the basis of total body weight and degree of vascular insufficiency.

The data set out in Table 1 was further analyzed to determine if the vascular status of the patients had any predictive value with respect to the efficacy of phentolamine mesylate in improving the erectile ability in impotent patients. The analysis showed that, of 16 total patients with normal vascular status, seven were successfully treated with phentolamine mesylate, six failed to respond, and three achieved partial erection. Of the 49 patients with vascular insufficiency, 15 were successful in achieving erection sufficient for vaginal penetration, while 29 failed. Five patients with vascular insufficiency achieved partial erection. The side effects seen in this study were infrequent and include, stuffy nose (6%), faintness or dizziness (2.3%) (relieved by lying down for 10-15 minutes) and vomiting 0.05%.

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### EXAMPLE 3

#### EFFECT OF BUCCAL PHENTOLAMINE MESYLATE AND IMPOTENCE OF VARIOUS ETIOLOGIES

Another single blind study was conducted in men having  
5     erectile dysfunction of various etiologies in order to further assess the efficacy  
of buccally administered phentolamine in ameliorating erectile dysfunction.

A mixed population of men having erectile dysfunction of  
various etiologies were given 3 filter paper strips impregnated with 20 mg of  
phentolamine mesylate and 3 placebo strips. The patients were not told which  
10    strips contained the drug and which strips were placebo. Patients were told  
to place one filter paper strip between the cheek and gum 10 minutes to 20  
minutes prior to attempts to achieve erection. The treatment was deemed  
successful if an erection sufficient to affect vaginal penetration was achieved.  
The results are shown in Table 2.

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TABLE 2

	Patient No.	Etiology	Result
	1	Peyronie's Disease and associated erectile dysfunction	2
	2	psychogenic	1
5	3	arteriogenic/psychogenic	1
	4	neurogenic	2
	5	arteriogenic	1
	6	arteriogenic	1
	7	*	2
10	8	*	2
	9	*	2
	10	*	2
	11	*	2
	12	psychogenic	No Report
15	13	arteriogenic	1
	14	psychogenic	1
	15	arteriogenic	2
	16	psychogenic	1
	17	psychogenic	1
20	18	arteriogenic/psychogenic	1
	19	arteriogenic	2
	20	psychogenic	1
	21	arteriogenic	2
	22	arteriogenic	2
25	23	arteriogenic/psychogenic	2

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TABLE 2 cont'd

- 5       \*       Prior to enrolling in this study, patients Nos. 7-11 were on a program of intracavernosal auto-injection using 20 $\mu$ g of PGE-1 with perfect efficacy. These patients also suffered from severe cardiovascular disease.

1 = erection and vaginal penetration

2 = failure to achieve erection sufficient for vaginal penetration

10       The results of this study establish that buccally administered phentolamine mesylate (20 mg) improved erectile ability in 36% of the patients reporting (excluding a total of two placebo responders). The results were most pronounced in patients with a diagnosis of psychogenic impotence wherein 5 of 5 patients reporting achieved an erection sufficient to vaginal penetration. Of those patients diagnosed as having arteriogenic impotence 3 of 7 patients showed improved erectile ability, with one of these patients also  
15       responding to placebo. Two of three patients diagnosed as having combined arteriogenic/psychogenic impotence showed improved erectile ability, with one of these patients also responding to placebo. Of the 5 patients in whom autoinjection of PGE-1 had shown perfect efficacy, none showed improvement in erectile ability with buccally administered phentolamine mesylate.

20       These results, when combined with those set out in Example 2, indicate that vasodilators such as phentolamine mesylate when administered via a route of administration which avoids the "first-pass" effect is effective in improving the onset of erectile ability in a significant percentage of impotent men and more particularly in men having psychogenic, arteriogenic,  
25       or combined psychogenic/arteriogenic impotence, the subpopulation of patients comprising the majority of cases of male impotence.

While useful in the treatment of impotence in males, the present invention is also useful in improving erectile ability in non-impotent men. It is well known that as men age, erectile ability may be altered. One

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manifestation of this decline in erectile ability is that the angle of erection subtended by the dorsal surface of the erect penis and the abdomen (erectile angles) increases with age. The method of the present invention, provides a means for improving erectile ability in non-impotent men by increasing blood  
5 flow to the penis, and thereby minimizing the erectile angle.

It should also be noted that filter strips impregnated with phentolamine mesylate lost their potency after several months of storage at room temperature in a paper bag.

#### EXAMPLE 4

#### 10 VASOACTIVE AGENTS USEFUL IN MODULATING THE HUMAN SEXUAL RESPONSE

A number of other vasoactive agents may be used in the practice of the present invention based on their demonstrated efficacy as vasodilators. Useful vasodilating drugs include those generally classified as  
15  $\alpha$ -adrenergic antagonists, sympathomimetic amines and those agents which exhibit direct relaxation of vascular smooth muscle.

Exemplary  $\alpha$ -adrenergic antagonists include phentolamine hydrochloride, phentolamine mesylate, phenoxybenzamine, tolazoline, dibenamine, yohimbine, and others. Phentolamine mesylate is preferred in the  
20 practice of the present invention. An exemplary sympathomimetic amine contemplated for use in the method of the present invention is nylidrin although other sympathomimetic amines having vasodilating activity are also comprehended by the invention.

Nicotinic acid (or nicotiny alcohol) has a direct vasodilating  
25 activity which is useful in the practice of the present invention. Papaverine is also non-specific smooth muscle relaxant which has vasodilating activity and has been used to treat male impotence by direct injection into the corpus

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cavernosum either alone or in combination with other drugs such as phentolamine.

Organic nitrates such as nitroglycerine and amyl nitrate also have pronounced vasodilating activity by virtue of their ability to relax  
5 vascular smooth muscle. Other vasoactive drugs of use in the practice of the present invention include but are not limited to thymoxamine, imipramine, verapamil, naftidrofuryl, isoxsuprine, and others.

In the practice of the present invention, these vasoactive agents are administered by the transmucosal, intranasal, transdermal, or rectal routes  
10 of administration such that the agent reaches its site of action prior to entering the portal circulation, thereby avoiding the "first-pass" effect.

Appropriate doses of each vasoactive agents for each route of administration are readily determined by those of ordinary skill in the art. By way of illustration, in order to determine the appropriate dose of each of the  
15 vasodilating agents of the present invention, one of ordinary skill in the art may use as a starting point, the usual published dosage of the vasodilator. The usual oral doses for commercially available vasodilators can be found in the Physician's Desk Reference published annually by Medical Economic Data, Montvale New Jersey, and in the available medical literature.

20 By way of example, Pavabid® oral papaverine hydrochloride is available from Marion Merrell Dow and is normally administered at 150 mg every 12 hours to achieve its vasodilating effects.

The oral dose of Calon® (verapamil hydrochloride) available from Searle is determined by titrating the individual patient with from 120 mg  
25 to about 240 mg of drug every 12 hours, the specific dose depending on the individual patient's response to the drug.

Yohimbine hydrochloride available as Daytohimbin® (Dayton Pharmaceuticals), Yocon (Palisades Pharmaceuticals), and Yohimex (Kramer) are all administered orally as 5.4 mg three times a day.

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Imipramine hydrochloride is available as Tofranil® from Geigy and is administered orally 4 times a day for a total dose ranging from 50 mg to about 150 mg per day.

5 Imipramine pamoate, also available from Geigy is administered in oral maintenance doses of 150 mg/day.

Using the established oral dosages as starting points, the optimal dosage for the specific route of administration can be determined by measuring baseline arterial blood flow in genital circulation of the patient prior to administration of the drug using a doppler ultrasound velocimeter as described in Example 1. Other methods such as thermography, plethysmography, radiometric or scintigraphic methods, and other methods well known in the art may also be utilized to assess blood flow in the genitalia. Having established base line blood flow, various dosages of the respective vasodilators may be administered by the routes of administration encompassed by the present invention and their effect on blood flow may be measured. The magnitude of the increase in blood flow necessary to modulate or enhance the sexual response in humans may vary from individual to individual, but is readily determined as described below. In addition, individual patients may be titrated with various dosages of the respective vasodilators until the optimum dosage is determined.

10  
15  
20

Vascular flow studies may also be coupled with assessments of sexual responsiveness as evidenced by the improvement of erectile ability in response to sexual stimulation.

### EXAMPLE 5

#### 25 MODULATION OF THE FEMALE SEXUAL RESPONSE

As discussed above, there are striking parallels between the vascular anatomy of male and female genitalia and in the erectile response facilitated by this vasculature. In both males and females, the erectile response

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takes place when under physical or psychological stimulation, blood flow to the genitalia increases by virtue of relaxation of smooth muscles in the arteries serving the genitalia.

5       The methods of the present invention may be used to improve or enhance the erectile response in women whose sexual response is impaired as evidenced by diminished capacity to produce sufficient vaginal lubrication to facilitate comfortable penile penetration and by other symptoms of impaired sexual responsiveness that may be correlated with the erectile response.

10       As in the case of male sexual response, in the absence of any clinically diagnosed dysfunction in the female erectile response, the methods of the present invention may be used to enhance the normal female sexual response. The "on demand" aspect of the present invention will allow a more rapid response to sexual stimulation along with heightened sensation associated with excitement and plateau stages of the female sexual response by virtue of  
15       the increased blood flow to the tissues.

In practice, enhancement of the female sexual response using the methods of the present invention are carried in much the same way as those described in Examples 2 and 3.

20       An effective vasodilating dose of a vasodilating agent is administered to a woman via the transmucosal, transdermal, intranasal, or rectal routes of administration. The appropriate doses of the particular vasodilating agent may be readily determined using methods described in Example 4. The female response may be measured using methods described in Masters, W.H. and Johnson, V.E., *Human Sexual Response*, Little, Brown,  
25       and Co., Boston (1966) which is incorporated herein by reference. Methods for measuring blood flow, including doppler ultrasonic velocimetry, thermography using for example an isothermal blood flow transducer, radiosciintigraphic methods, photoplethysmography may be used as well as other methods well known in the art. In addition, measuring the contraction  
30       of the distal 1/3 as is characteristic of the plateau phase of female sexual

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response of the vagina may be measured using methods and equipment well known in the art including but not limited to strain gauges or other devices for measuring muscular contraction or muscle tension.

5 In addition, enhanced sexual response may be measured in a more subjective manner by simply asking the female subject to describe any change in sensation brought about by administration of the vasodilator by the methods of the present invention. Appropriate placebo controls should also be conducted to ascertain whether or not the effort is directly attributable to the administration of the vasodilator.

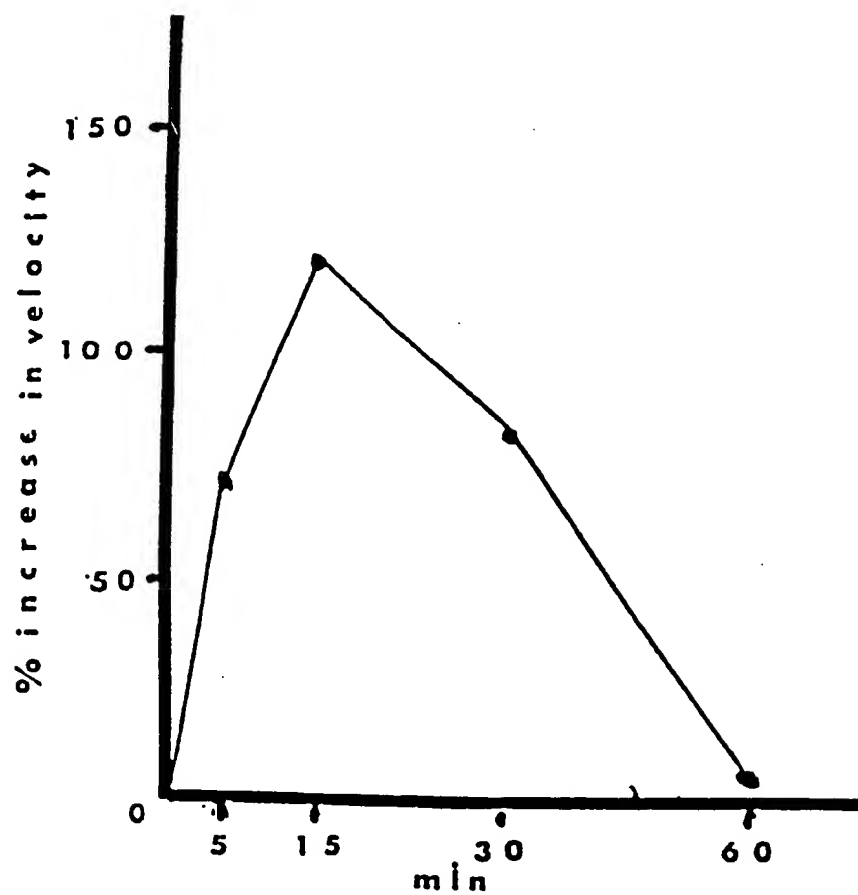
10 A preferred embodiment of the present invention involves the buccal administration of from about 5 mg to about 80 mg of phentolamine mesylate from about 1 minute to about 1 hour prior to, and in preparation for, intercourse. However any of the other routes of administration of any of the vasodilating agent included within the scope of the present invention may be  
15 used.

While this invention has been described by way of preferred embodiments, the examples set out herein are not intended to limit the scope of the invention which contemplates the use of any pharmacologic vasodilating drug capable of absorption into the systemic circulation upon administration  
20 of the drug via the transmucosal, transdermal, intranasal, or rectal routes of administration.

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PCT CLAIMS

We claim:

1. The use of a drug having vasodilator activity for the manufacture of a medicament for transmucosal, transdermal, intranasal, and rectal administration to modulate sexual response in a human.
- 5                   2. The use of claim 1 wherein the drug is selected from the group consisting of phentolamine mesylate, phentolamine hydrochloride, phenoxybenzamine, yohimbine, nitroglycerine, thymoxamine, imipramine, verapamil, isoxsuprine, naftidrofuryl, tolazoline, and papaverine.
- 10                   3. The use of a drug comprising phentolamine mesylate for the manufacture of a medicament for transmucosal administration to treat male impotence.
4. The use according to claim 3 wherein the medicament is in tablet form.
- 15                   5. The use according to claim 4, wherein each tablet contains about 5 mg to about 80 mg of phentolamine mesylate.

**FIGURE 1**

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US94/09048**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(5) : Please See Extra Sheet.

US CL : 514/401, 651, 280, 509, 546, 277, 217, 654, 471, 307

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/401, 651, 280, 509, 546, 277, 217, 654, 471, 307

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

STN (REGISTRY, CHEMICAL ABSTRACTS, BIOSIS, MEDLINE), APS

search terms: names of compounds in claim 2, sexual response, sexual dysfunction, impotence, erection

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X -- Y	US, A, 4,801,587 (VOSS ET AL) 31 JANUARY 1989, see the abstract and the claims.	1-3 ----- 4-5
X -- Y	US, A, 5,059,603 (RUBIN) 22 OCTOBER 1991, see particularly column 4, line 30-51.	1-3 ----- 4-5
Y	US, A, 3,943,246 (STURMER) 09 MARCH 1976, see entire document.	4-5

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier document published on or after the international filing date	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&" document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means	
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search 05 OCTOBER 1994	Date of mailing of the international search report 01 NOV 1994
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230	Authorized Officer W. JARVIS & K. MACMILLAN Telephone No. (703) 308-1235

**INTERNATIONAL SEARCH REPORT**

International application No.  
PCT/US94/09048

**A. CLASSIFICATION OF SUBJECT MATTER:**

IPC (5):

A61K 31/415, 31/135, 31/44, 31/21, 31/22, 31/435, 31/55, 31/34, 31/47